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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/625,471	07/23/2003	Angel Pellicer	PELLICERIA	1686

7590 08/02/2007  
BROWDY AND NEIMARK, P.L.L.C.  
624 Ninth Street, N.W.  
Washington, DC 20001-5303

EXAMINER
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HOLLERAN, ANNE L

ART UNIT	PAPER NUMBER
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1643

MAIL DATE	DELIVERY MODE
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08/02/2007

PAPER

**Please find below and/or attached an Office communication concerning this application or proceeding.**

The time period for reply, if any, is set in the attached communication.

<b>Office Action Summary</b>	<b>Application No.</b> 10/625,471	<b>Applicant(s)</b> PELLICER ET AL.	
	<b>Examiner</b> Anne L. Holleran	<b>Art Unit</b> 1643	

**-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --**

**Period for Reply**

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

**Status**

- 1) ☒ Responsive to communication(s) filed on 14 December 2006.
- 2a) ☒ This action is **FINAL**.                      2b) ☐ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

**Disposition of Claims**

- 4) ☒ Claim(s) 16-18, 20, 21, 23-35 is/are pending in the application.
- 4a) Of the above claim(s) 27-33 is/are withdrawn from consideration.
- 5) ☐ Claim(s) \_\_\_\_\_ is/are allowed.
- 6) ☒ Claim(s) 16, 20, 21, 23, 24, 26, 34 and 35 is/are rejected.
- 7) ☒ Claim(s) 17, 18 and 25 is/are objected to.
- 8) ☐ Claim(s) \_\_\_\_\_ are subject to restriction and/or election requirement.

**Application Papers**

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on \_\_\_\_\_ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.  
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).  
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
- 11) ☐ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

**Priority under 35 U.S.C. § 119**

- 12) ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☐ All    b) ☐ Some \* c) ☐ None of:
1. ☐ Certified copies of the priority documents have been received.
  2. ☐ Certified copies of the priority documents have been received in Application No. \_\_\_\_\_.
  3. ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

\* See the attached detailed Office action for a list of the certified copies not received.

**Attachment(s)**

- |  |   |
|--|---|
| 1) <input checked="" type="checkbox"/> Notice of References Cited (PTO-892)                                | 4) <input type="checkbox"/> Interview Summary (PTO-413)<br>Paper No(s)/Mail Date. _____ |
| 2) <input type="checkbox"/> Notice of Draftsperson's Patent Drawing Review (PTO-948)                       | 5) <input type="checkbox"/> Notice of Informal Patent Application                       |
| 3) <input type="checkbox"/> Information Disclosure Statement(s) (PTO/SB/08)<br>Paper No(s)/Mail Date _____ | 6) <input type="checkbox"/> Other: _____  |

**DETAILED ACTION**

1. This Office action supersedes the previous Office action, mailed 3/28/2007. The previous Office action is vacated.
2. The amendment file 12/14/2006 is acknowledged. Claims 1-15, 19 and 22 were canceled.
3. Claims 16-18, 20, 21, 23-35 are pending. Claims 27-33, drawn to non-elected inventions, are withdrawn from consideration.

Claims 16-18, 20, 21, 23-26, 34 and 35 are examined on the merits.

***Claim Objections/Rejections Withdrawn:***

4. The objection to claims 16, 19 and 20 for depending from a claim that is withdrawn from consideration is withdrawn in view of the amendment to claim 16 and 20, and the cancellation of claim 19.

***Claim Rejections - 35 USC § 112***

5. The rejection of claims 16, 20, 34 and 35 under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention is withdrawn in view of the amendment to the claims.

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6. The rejection of claims 16, 20, 34 and 35 under 35 U.S.C. 102(b) as being anticipated by D'Adamo (D'Adamo, et al., Oncogene, 14: 1295-1305, 1997; cited in the IDS) or Miller (Miller, M.J. et al., Journal of Biological Chemistry, 272(9): 5600-5605, 1997) is withdrawn in view of the amendments to the claims.

***Claim Rejections Maintained and New Grounds of Rejection:***

***Claim Rejections - 35 USC § 112***

The following is a quotation of the second paragraph of 35 U.S.C. 112:

The specification shall conclude with one or more claims particularly pointing out and distinctly claiming the subject matter which the applicant regards as his invention.

7. Claims 23 and 26 are rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.

Claim 23 (and dependent claim 26) is indefinite because it appears to be missing a phrase such as, for example, "the abnormally truncated variant".

The following is a quotation of the first paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

8. Claims 16, 20, 21, 34 and 35 remain rejected under 35 U.S.C. 112, first paragraph, because the specification, while being enabling for isolated nucleic acids that encode a polypeptide that comprises the amino acid sequence of SEQ ID NO: 2, or for nucleic acids *consisting of* nucleic acid sequences that encode SEQ ID NO: 8, or for specific nucleic acids

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*consisting of* the sequences of SEQ ID NOS: 5-7 and 9, 11, 13, 15, 17, 19, 21 and 23, does not reasonably provide enablement for nucleic acids encoding a polypeptide that is a variant of human Rgr consisting of an amino acid sequence with at least 98% sequence identity to SEQ ID NO: 2, or *comprises* the nucleic acid sequences of SEQ ID NOS: 5-7. The specification does not enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the invention commensurate in scope with these claims.

Applicants argue that the amendments to the claims obviate the rejection of record. This is not found persuasive because, as explained in the previous Office action, it is not clear from the specification how one of skill in the art may use nucleic acids that encode polypeptides having functional activity that is not the same as the amino acid sequence of SEQ ID NO: 2, which encodes human Rgr. In the previous Office action, the examiner noted that even a single amino acid substitution will often dramatically affect the biological activity and characteristics of a protein, and that the specification does not teach specific biological activities of human Rgr, and that it is unpredictable whether human Rgr exhibits exactly the same set of activities as rabbit Rgr. Therefore, the claims to nucleic acids encoding variant polypeptides of human Rgr consisting of an amino acid sequence with at least 98% sequence identity to the amino acid sequence of SEQ ID NO: 2 are not enabled by the specification because one of skill in the art would first have to establish what are the activities of human Rgr, and then to screen for the working embodiments. Without a set of activities to use in a routine assay, it would require further and undue experimentation to screen for useful variants of human Rgr protein consisting of an amino acid sequence with at least 98% sequence identity to SEQ ID NO: 2. Furthermore, the claim 16 and also claim 20 contains the recitation “consisting of an amino acid sequence with

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at least 98% identity to SEQ ID NO: 2” reads on subsequences of SEQ ID NO: 2, where the amino acid sequences of the subsequences has 98% sequence identity with SEQ ID NO: 2.

Applicants also explain that in one aspect the claims are limited to polynucleotides consisting of the nucleic acid sequence of SEQ ID NO: 5, SEQ ID NO: 6 or SEQ ID NO: 7. However, in claim 16 the claims are drawn to “isolated nuclei acid molecules comprising a nucleotide sequence...”. Therefore, the claims read on nucleic acid sequences comprising fragments of a polynucleotide encoding SEQ ID NO: 2.

9. Claims 16, 20 and 21 remain rejected under 35 U.S.C. 112, first paragraph, as failing to comply with the written description requirement. The claim(s) contains subject matter that was not described in the specification in such a way as to reasonably convey to one skilled in the relevant art that the inventor(s), at the time the application was filed, had possession of the claimed invention. The basis for this rejection is that the specification fails to provide a description that is sufficient to support genus claims.

Applicants assert that the claim amendments obviate the rejections under 35 USC 112, first paragraph. This is not found persuasive because claim 16 continues to be drawn to “nucleic acid molecules comprising a nucleotide sequence”, and because claim 20 recites “wherein the polypeptide consists of an amino acid sequence with at least 98% sequence identity to SEQ ID NO: 2, which reads on subsequences. Claim 21 depends from claim 20, and appears to read on subsequences because it is dependent from claim 20. Therefore, given the preamble of claim 16, and because claims 20 and 21 depend from claim 16, claims 20 and 21 appear to be drawn to polynucleotides comprising fragments of polynucleotides encoding SEQ ID NO: 2. In the

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previous Office action it was pointed out that the specification only teaches fragments of SEQ ID NO: 2 in the context of SEQ ID NO: 2 itself, or in the context of the specific fragment itself (consisting of the fragment).

***Claim Rejections - 35 USC § 102***

The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless –

(b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.

(e) the invention was described in (1) an application for patent, published under section 122(b), by another filed in the United States before the invention by the applicant for patent or (2) a patent granted on an application for patent by another filed in the United States before the invention by the applicant for patent, except that an international application filed under the treaty defined in section 351(a) shall have the effects for purposes of this subsection of an application filed in the United States only if the international application designated the United States and was published under Article 21(2) of such treaty in the English language.

(a) the invention was known or used by others in this country, or patented or described in a printed publication in this or a foreign country, before the invention thereof by the applicant for a patent.

10. Claims 16 remains rejected under 35 U.S.C. 102(e) as being anticipated by WO 01/57278 (Penn et al., 9 August 2001).

Applicants argue that the rejection is obviated by the amendments, because the claims no longer read on polynucleotides comprising nucleic acids comprising the sequence of SEQ ID NO: 5, 6 or 7. This is not found persuasive because the preamble of claim 16 is drawn to “an isolated nucleic acid molecule comprising a nucleotide sequence...”

Penn teaches nucleic acid molecules that comprise SEQ ID NO: 5, 6 or 7. Therefore, Penn teaches the claimed nucleic acid molecules.

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11. Claims 16, 20 and 21 are rejected under 35 U.S.C. 102(b) as being anticipated by Accession No. AAA97456 (CN1257923-A in database Geneseq, Gencore version 5.1.9, 28 June 2000).

Claims 16 and 20 are drawn to nucleic acid molecules where the polypeptide consists of an amino acid sequence with at least 98% sequence identity to SEQ ID NO: 2. Because of the phrase “consisting of an amino acid sequence”, the claims read on nucleotides encoding subsequences of SEQ ID NO: 2, and because of the preamble of claim 16, where claim 16 is drawn to “an isolated nucleic acid molecule comprising a nucleotide sequence...”, claims 16, 20 and 21 are drawn to sequences comprising polynucleotides encoding subsequences of SEQ ID NO: 2.

Accession No. AAA97456 comprises subsequences of SEQ ID NO: 2 (see alignment) and therefore teaches polynucleotides within the scope of the claims.

12. Claims 16, 20 and 21 are rejected under 35 U.S.C. 102(a) as being anticipated by Accession No. AAS34856 (WO200155163-A in database Geneseq, Gencore version 5.1.9, 02 August 2001).

Claims 16 and 20 are drawn to nucleic acid molecules encoding polypeptides, where the polypeptides consist of an amino acid sequence with at least 98% sequence identity to SEQ ID NO: 2. Because of the phrase “consisting of an amino acid sequence”, the claims read on nucleotides encoding subsequences of SEQ ID NO: 2, and because of the preamble of claim 16, where claim 16 is drawn to “an isolated nucleic acid molecule comprising a nucleotide



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sequence...”, claims 16, 20 and 21 are drawn to sequences comprising polynucleotides encoding subsequences of SEQ ID NO: 2.

Accession No. AAS34856 comprises subsequences of SEQ ID NO: 2 (see alignment) and therefore teaches polynucleotides within the scope of the claims.

13. Claims 16, 20, 21, 23, 34 and 35 are rejected under 35 U.S.C. 102(e) as being anticipated by Tang (WO 01/53312 A1; publication date 26 July 2001; international filing date is 26 December 2000; pages 2-5, 27-32, 299, and 6443).

Tang discloses SEQ ID NO: 3567 (see page 6443) which is an amino acid sequence that encompasses SEQ ID NOs: 10 and 22. Tang discloses polynucleotides encoding SEQ ID NO: 3567 (see pages 2-5, 27 and 299), which is a sequence that encodes SEQ ID NO: 10, 22, and encodes a sequence that is at least 98% identical to SEQ ID NO: 2. Tang discloses methods of making polypeptides that require the use of host cells transfected with vectors comprising the encoding polynucleotides (see page 4, lines 4-10; page 29, lines 19-27). The claims are drawn to nucleic acids comprising polynucleotides encoding polypeptides consisting of amino acid sequences such as SEQ ID NO: 10 and 22, vectors and host cells. Even though the claims contain the transitional phrase “consisting of” with respect to the polypeptide sequence, the claims are drawn to nucleic acid molecules that comprise polynucleotide sequences encoding those polypeptide sequences. Therefore, Tang teaches polynucleotides, vectors and host cells that are the same as that claimed.

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14. Claims 16, and 23 are rejected under 35 U.S.C. 102(b) as being anticipated by either Adams (Accession No. AA311687, in Nature 377 (6547 Suppl): 3-174, 1995) or Hedge (Accession No. AW962844, 01 June 2000).

Claims 16 and 23 read on isolated nucleic acid molecules comprising a nucleotide sequence encoding for a polypeptide selected from the group of consisting of SEQ ID NO: 10, 12, 14, 16, 18, 20, 22 or 24.

Either of Adams or Hedge teaches nucleic acids that encode SEQ ID NO: 22 (see following alignments for SEQ ID NO: 22). Adams or Hedge also teach nucleic acids that encode SEQ ID NO: 10. Therefore, either of Adams or Hedge teach nucleic acid molecules that are the same as that claimed.

```

RESULT 1
AA311687
LOCUS      AA311687          408 bp    mRNA    linear    EST 19-APR-1997
DEFINITION EST182411 Jurkat T-cells VI Homo sapiens cDNA 5' end similar to
            similar to guanine nucleotide dissociation stimulator, mRNA
            sequence.
ACCESSION  AA311687
VERSION    AA311687.1  GI:1964015
KEYWORDS   EST.
SOURCE     Homo sapiens (human)
            ORGANISM
            Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Euarchontoglires; Primates; Haplorrhini;
            Catarrhini; Hominidae; Homo.
REFERENCE  1  (bases 1 to 408)
AUTHORS    Adams,M.D., Kerlavage,A.R., Fleischmann,R.D., Fuldner,R.A.,
            Bult,C.J., Lee,N.H., Kirkness,E.F., Weinstock,K.G., Gocayne,J.D.,
            White,O., Sutton,G., Blake,J.A., Brandon,R.C., Man-Wai,C.,
            Clayton,R.A., Cline,T.R., Cotton,M.D., Earle-Hughes,J., Fine,L.D.,
            Fitzgerald,L.M., Fitzhugh,W.M., Fritchman,J.L., Geoghagen,N.S.,
            Glodek,A., Gnehm,C.L., Hanna,M.C., Hedblom,E., Hinkle,P.S.Jr.,
            Kelley,J.M., Kelley,J.C., Liu,L.-I., Marmaros,S.M., Merrick,J.M.,
            Moreno-Palanques,R.F., McDonald,L.A., Nguyen,D.T., Pelligrino,S.M.,
            Phillips,C.A., Ryder,S.E., Scott,J.L., Saudek,D.M., Shirley,R.,
            Small,K.V., Spriggs,T.A., Utterback,T.R., Weidman,J.F., Li,Y.,
            Bednarik,D.P., Cao,L., Cepeda,M.A., Coleman,T.A., Collins,E.J.,
            Dimke,D., Feng,D.-F., Ferrie,A., Fischer,C., Hastings,G.A.,
            He,W.W., Hu,J.S., Greene,J.M., Gruber,J., Hudson,P., Kim,A.K.,
            Kozak,D.L., Kunsch,C., Hungjun,J., Li,H., Meissner,P.S., Olsen,H.,
            Raymond,L., Wei,Y.F., Wing,J., Xu,C., Yu,G.L., Ruben,S.M.,
            Dillion,P.J., Fannon,M.R., Rosen,C.A., Haseltine,W.A., Fields,C.,
            Fraser,C.M. and Venter,J.C.
TITLE      Initial assessment of human gene diversity and expression patterns
            based upon 83 million nucleotides of cDNA sequence
JOURNAL    Nature 377 (6547 Suppl), 3-174 (1995)
PUBMED     7566098
COMMENT    Other_ESTs: THC123926

```

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Contact: Kerlavage, AR  
 Bioinformatics  
 The Institute for Genomic Research  
 9712 Medical Center Drive, Rockville, MD 20850 USA  
 Tel: 3018699056  
 Fax: 3018699423  
 Email: arkerlav@tigr.org  
 For clone availability, additional sequence and expression  
 information related to this EST, please check the TIGR Human Gene  
 Index (<http://www.tigr.org/tdb/hgi/hgi.html>)  
 Seq primer: M13 Reverse.

FEATURES  
 source Location/Qualifiers  
 1..408  
 /organism="Homo sapiens"  
 /mol\_type="mRNA"  
 /db\_xref="ATCC (inhost):158906"  
 /db\_xref="taxon:9606"  
 /cell\_type="T-lymphocyte"  
 /clone\_lib="Jurkat T-cells VI"  
 /note="Vector: pBluescript SK-; Site\_1: EcoRI; Site\_2:  
 XhoI"

## ORIGIN

## Alignment Scores:

Pred. No.:	1.18e-56	Length:	408
Score:	478.00	Matches:	94
Percent Similarity:	100.0%	Conservative:	0
Best Local Similarity:	100.0%	Mismatches:	0
Query Match:	100.0%	Indels:	0
DB:	1	Gaps:	0

US-10-625-471-22 (1-94) x AA311687 (1-408)

```

Qy      1 MetArgLeuArgArgGlnLysLysGlyValValProPheLeuGlyAspPheLeuThrGlu 20
      |||
Db      16 ATGAGGCTGCGGAGGCAGAAGAAGGGTGTGGTCCCTTCCTGGGGGATTTCTGACTGAG 75

Qy      21 LeuGlnArgLeuAspSerAlaIleProAspAspLeuAspGlyAsnThrAsnLysArgSer 40
      |||
Db      76 TTACAGAGGCTGGATTTCGGCCATCCCGGACGACCTGGATGGCAACACCAACAAGAGGAGC 135

Qy      41 LysGluValArgValLeuGlnGluMetGlnLeuLeuGlnValAlaAlaMetAsnTyrArg 60
      |||
Db      136 AAGGAGGTCCGAGTTCTGCAGGAAATGCAGCTGCTCCAAGTGGCTGCCATGAATTACAGG 195

Qy      61 LeuArgProLeuGluLysPheValThrTyrPheThrArgMetGluGlnLeuSerAspLys 80
      |||
Db      196 CTCGGCCTCTTGAGAAATTTGTACCTATTTTACAAGAATGGAGCAGCTCAGTGACAAA 255

Qy      81 GluSerTyrLysLeuSerCysGlnLeuGluProGluAsnPro 94
      |||
Db      256 GAGAGCTACAAGCTNTCTGCCAGCTGGAGCCCGAAAACCCG 297

```

## RESULT 2

AW962844  
 LOCUS AW962844 583 bp mRNA linear EST 01-JUN-2000  
 DEFINITION EST374917 MAGE resequences, MAGG Homo sapiens cDNA, mRNA sequence.  
 ACCESSION AW962844  
 VERSION AW962844.1 GI:8152680  
 KEYWORDS EST.  
 SOURCE Homo sapiens (human)  
 ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Euarchontoglires; Primates; Haplorrhini;  
 Catarrhini; Hominidae; Homo.  
 REFERENCE 1 (bases 1 to 583)  
 AUTHORS Hegde, P., Qi, R., Abernathy, K., Dharap, S., Gaspard, R., Gay, C.,

	Holt, I.E., Saeed, A.I., Sharov, V., Lee, N.H., Yeatman, T.J. and Quackenbush, J.
TITLE	Assessment of gene expression patterns in a model of colon tumor metastasis using a 19,200 element cDNA microarray
JOURNAL	Unpublished (2000)
COMMENT	Contact: John Quackenbush The Institute for Genomic Research 9712 Medical Center Dr., Rockville, MD 20850, USA Tel: 301 838 3528 Fax: 301 838 0208 Email: johnq@tigr.org Plate: 180 Seq primer: Reverse.

```
FEATURES
    source          1. .583
                   /organism="Homo sapiens"
                   /mol_type="mRNA"
                   /db_xref="taxon:9606"
                   /clone_lib="MAGE resequences, MAGG"
                   /vector="Vector: pBluescriptSKm"
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ORIGIN

Alignment Scores:

Pred. No.:	1.88e-56	Length:	583
Score:	478.00	Matches:	94
Percent Similarity:	100.0%	Conservative:	0
Best Local Similarity:	100.0%	Mismatches:	0
Query Match:	100.0%	Indels:	0
DB:	7	Gaps:	0

US-10-625-471-22 (1-94) x AW962844 (1-583)

Qy	1	MetArgLeuArgArgGlnLysLysGlyValValProPheLeuGlyAspPheLeuThrGlu	20
Db	15	ATGAGGCTGCGGAGGCAGAGAAGGGTGTGGTCCCCTTCTGGGGGATTTTCTGACTGAG	74
Qy	21	LeuGlnArgLeuAspSerAlaIleProAspAspLeuAspGlyAsnThrAsnLysArgSer	40
Db	75	TTACAGAGGCTGGATTCGGCCATCCCGACGACCTGGATGGCAACACCAACAAGAGGAGC	134
Qy	41	LysGluValArgValLeuGlnGluMetGlnLeuLeuGlnValAlaAlaMetAsnTyrArg	60
Db	135	AAGGAGGTCCGAGTTCTGCAGGAAATGCAGCTGCTCCAAGTGGCTGCCATGAATTACAGG	194
Qy	61	LeuArgProLeuGluLysPheValThrTyrPheThrArgMetGluGlnLeuSerAspLys	80
Db	195	CTTCGGCCTCTTGAGAAATTGTACCTATTTACAAGAATGGAGCAGCTCAGTGACAAA	254
Qy	81	GluSerTyrLysLeuSerCysGlnLeuGluProGluAsnPro	94
Db	255	GAGAGCTACAAGCTGTCTGCCAGCTGGAGCCCCGAAAACCCG	296

15. Claims 16 and 23 remain rejected under 35 U.S.C. 102(a) as being anticipated by

Accession No. BI837800, (NIH-Mammalian Gene Collection, 04 Oct. 2001).

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Claims 16 and 23 read on isolated nucleic acid molecules comprising a nucleotide sequence encoding for a polypeptide selected from the group of consisting of SEQ ID NO: 10, 12, 14, 16, 18, 20, 22 or 24.

NIH-MGC teaches nucleic acids that encode SEQ ID NO: 20 (see following alignments for SEQ ID NO: 20). NIH-MGC also teaches a nucleic acid that encodes SEQ ID NO: 14.

Therefore, NIH-MGC teaches nucleic acid molecules that are the same as that claimed.

```

RESULT 1
BI837800
LOCUS      BI837800              707 bp    mRNA    linear    EST 04-OCT-2001
DEFINITION 603083504F1 NIH_MGC_120 Homo sapiens cDNA clone IMAGE:5222808 5',
            mRNA sequence.
ACCESSION  BI837800
VERSION    BI837800.1  GI:15949350
KEYWORDS   EST.
SOURCE     Homo sapiens (human)
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Euarchontoglires; Primates; Haplorrhini;
            Catarrhini; Hominidae; Homo.
REFERENCE  1 (bases 1 to 707)
AUTHORS    NIH-MGC http://mgc.nci.nih.gov/.
TITLE      National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL    Unpublished (1999)
COMMENT     Contact: Robert Strausberg, Ph.D.
            Email: cgapbs-r@mail.nih.gov
            Tissue Procurement: Life Technologies, Inc.
            cDNA Library Preparation: Life Technologies, Inc.
            cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
            DNA Sequencing by: Incyte Genomics, Inc.
            Clone distribution: MGC clone distribution information can be
            found through the I.M.A.G.E. Consortium/LLNL at:
            http://image.llnl.gov
            Plate: LLAM11560 row: g column: 01
            High quality sequence stop: 707.
FEATURES   Location/Qualifiers
            source
                1..707
                /organism="Homo sapiens"
                /mol_type="mRNA"
                /db_xref="taxon:9606"
                /clone="IMAGE:5222808"
                /lab_host="DH10B"
                /clone_lib="NIH_MGC_120"
                /note="Organ: pooled pancreas and spleen; Vector:
                pCMV-SPORT6; Site_1: NotI; Site_2: EcoRV (destroyed); RNA
                source anonymous pool of spleen and pancreas from 28 yo
                male. Library is oligo-dT primed and directionally cloned
                (EcoRV site is destroyed upon cloning). Average insert
                size 1.5 kb, insert size range 1-2.5 kb. Library is
                normalized and enriched for full-length clones and was
                constructed by C. Gruber (Invitrogen). Research Genetics
                tracking code 025. Note: this is a NIH_MGC Library."
ORIGIN
Alignment Scores:
Pred. No.:      1.03e-78      Length:      707

```

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Score:	773.00	Matches:	150
Percent Similarity:	100.0%	Conservative:	0
Best Local Similarity:	100.0%	Mismatches:	0
Query Match:	100.0%	Indels:	0
DB:	3	Gaps:	0

US-10-625-471-20 (1-150) x BI837800 (1-707)

```

Qy      1 MetSerThrValProGlyGlySerArgHisSerLeuGlyIleGlnValArgGlyGlyTrp 20
      |||
Db      14 ATGAGCACGGTGCCAGGTGGCTCCCGCCACTCCCTGGGGATCCAAGTGC GG GTGGCTGG 73

Qy     21 GlyValThrGlyGlyGluGluGluSerLeuThrValProValAlaAspThrTrpGlnAla 40
      |||
Db      74 GGTGTAACTGGGGGAGAGGAGAGAGAGCCTCACTGTCCCTGTCGCTGACACCTGGCAGGCG 133

Qy     41 GlySerPheLysValAlaThrGlnGluArgAsnProGlnArgValGlnMetArgLeuArg 60
      |||
Db     134 GGGAGCTTTAAGGTGGCCACCCAGGAGAGGAACCCCGAGAGAGTCCAGATGAGGCTGCGG 193

Qy     61 ArgGlnLysLysGlyValValProPheLeuGlyAspPheLeuThrGluLeuGlnArgLeu 80
      |||
Db     194 AGGCAGAAGAAGGGTGTGGTCCCTTCCTGGGGGATTTTCTGACTGAGTTACAGAGGCTG 253

Qy     81 AspSerAlaIleProAspAspLeuAspGlyAsnThrAsnLysArgSerLysGluValArg 100
      |||
Db     254 GATTGCGCCATCCCGACGACCTGGATGGCAACACCAACAAGAGGAGCAAGGAGGTCCGA 313

Qy    101 ValLeuGlnGluMetGlnLeuLeuGlnValAlaAlaMetAsnTyrArgLeuArgProLeu 120
      |||
Db     314 GTTCTGCAGGAAATGCAGCTGTCTCAAGTGGCTGCCATGAATTACAGGCTTCGGCCTCTT 373

Qy    121 GluLysPheValThrTyrPheThrArgMetGluGlnLeuSerAspLysGluSerTyrLys 140
      |||
Db     374 GAGAAATTTGTACCTATTTTCAAGAATGGAGCAGCTCAGTGACAAAGAGAGCTACAAG 433

Qy    141 LeuSerCysGlnLeuGluProGluAsnPro 150
      |||
Db     434 CTGTCTTGCCAGCTGGAGCCCGAAAACCCG 463

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### *Conclusion*

No claim is allowed. Claims 17, 18 and 25 are objected for depending from a rejected claim.

Applicant's amendment necessitated the new ground(s) of rejection presented in this Office action. Accordingly, **THIS ACTION IS MADE FINAL**. See MPEP § 706.07(a). Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after

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the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the date of this final action.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Anne Holleran, whose telephone number is (571) 272-0833. The examiner can normally be reached on Monday through Friday from 9:30 am to 5:00 pm. If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Larry Helms, can be reached on (571) 272-0832. Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the Group receptionist whose telephone number is (571) 272-1600.

Papers related to this application may be submitted to Group 1600 by facsimile transmission. The faxing of such papers must conform to the notice published in the Official Gazette, 1096 OG 30 (November 15, 1989). The Official Fax number for Group 1600 is (571) 273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR

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system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private

PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll free).

Anne L. Holleran

Patent Examiner

March 7, 2007

  
ALANA M. HARRIS, PH.D.  
PRIMARY EXAMINER